Reshma KOTIAN et al., Serial No. 10/768,348 "Stabilized Paroxetine Formulation"

IN THE CLAIMS

- 26. (Amended) A moisture barrier, non-controlled release pharmaceutical excipient solution comprising:
 - A) ethylcellulose;

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- B) polar solvent;
- C) alcohol; and
- D) a surfactant.
- 27 (Original) The moisture barrier pharmaceutical excipient of claim 26, wherein said surfactant comprises polysorbate 80, and wherein said polysorbate 80 and said ethylcellulose are present in a ratio of approximately 1.00: 0.165.
- 28. (Amended) A process for manufacturing a substantially moisture stable, non-controlled release pharmaceutical product, comprising:
 - A) mixing ethylcellulose, polar solvent, alcohol and a surfactant to make a moisture barrier, non-controlled release pharmaceutical excipient solution;
 - B) mixing a drug substance with said moisture barrier pharmaceutical excipient solution to form substantially moisture stable drug substance;
 - C) coating the substantially moisture-resistant drug substance of step B in a pharmaceutically-acceptable coating.
- 29. (Original) The process of claim 28, wherein said pharmaceutically-acceptable coating comprises a gelatin capsule.

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30. (Original) The process of claim 28, wherein said surfactant comprises polysorbate 80. and wherein said polysorbate 80 and said ethylcellulose are present in a ratio of approximately 1.00:0.165.

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- 31. (Original) The process of claim 28, wherein said drug substance comprises paroxetine.
- (Original) The process of claim 3, wherein said surfactant comprises polysorbate 80, and wherein said polysorbate 80 and said ethylcellulose are present in a ratio of approximately 1.00: 0.165, and wherein said drug substance comprises paroxetine.
- 33. (Amended) A substantially moisture stable, non-controlled release drug product comprising:
 - A) A substantially moisture stable, non-controlled release core comprising a drug substance, ethylcellulose and a surfactant; and
 - B) An outer layer surrounding said core, said outer layer comprising a pharmaceutically acceptable material, said outer layer substantially free of said drug substance.
- 34. (Original) The drug product of claim 33, wherein:
 - A) said surfactant comprises polysorbate 80, and wherein said polysorbate 80 and said ethylcellulose are present in a ratio of approximately 1.00: 0.165.
- 35. (Original) The drug product of claim 33, wherein:
 - A) said drug substance comprises paroxetine.
- 36. (Original) The drug product of claim 33, wherein:
 - A) said drug substance comprises paroxetine, said surfactant comprises polysorbate 80, and wherein said polysorbate 80 and said ethylcellulose are present in a ratio of approximately 1.00: 0.165.

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